

Claims

1. A method for producing a transgenic plant comprising:
- 5 i) transforming a plant cell with a first expression cassette comprising a nucleic acid sequence encoding a D-amino acid oxidase operably linked with a promoter allowing expression in plant cells or plants, in combination with at least one second expression cassette suitable for conferring to said plant an agronomically valuable trait, and
- 10 ii) providing at least one first compound X, which is phytotoxic against plant cells not functionally expressing said D-amino acid oxidase, wherein said compound X can be metabolized by said D-amino acid oxidase into one or more compound(s) Y which are non-phytotoxic or less phytotoxic than compound X, and
- 15 iii) treating said transformed plant cells of step i) with said first compound X in a phytotoxic concentration and selecting plant cells comprising in their genome both said first and said second expression cassette, wherein said first expression cassette is conferring resistance to said transformed plant cells against said compound X by expression of said D-amino acid oxidase, and
- 20 iv) providing at least one second compound M, which is non-phytotoxic or moderately phytotoxic against plant cells not functionally expressing said D-amino acid oxidase, wherein said compound M can be metabolized by said D-amino acid oxidase into one or more compound(s) N which are phytotoxic or more phytotoxic than compound M, and
- 25 v) breaking the combination between said first expression cassette and said second expression cassette and treating resulting said plant cells with said second compound M in a concentration toxic to plant cells still comprising said first expression cassette, and selecting plant cells comprising said second expression cassette but lacking said first expression cassette.
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- 35 2. The method of Claim 1, wherein said first expression cassette for said D-amino acid oxidase and said second expression cassette for said agronomically valuable trait are

- a) both comprised in one DNA construct and combination is broken by deletion or excision of said first expression cassette for said D-amino acid oxidase, or
- 5 b) are comprised on separate DNA constructs which are transformed in combination by co-transformation into said plant cells, and combination is broken by subsequent segregation of the two expression cassettes.
- 10 3. The method of Claim 1 and 2, wherein said method for producing a transgenic plant comprises the steps of:
- i) transforming a plant cell with a first DNA construct comprising
- 15 a) a first expression cassette comprising a nucleic acid sequence encoding a D-amino acid oxidase operably linked with a promoter allowing expression in plant cells or plants, wherein said first expression cassette is flanked by sequences which allow for specific deletion of said first expression cassette, and
- 20 b) at least one second expression cassette suitable for conferring to said plant an agronomically valuable trait, wherein said second expression cassette is not localized between said sequences which allow for specific deletion of said first expression cassette, and
- 25 ii) providing at least one first compound X, which is phytotoxic against plant cells not functionally expressing said D-amino acid oxidase, wherein said compound X can be metabolized by said D-amino acid oxidase into one or more compound(s) Y which are non-phytotoxic or less phytotoxic than compound X, and
- 30 iii) treating said transformed plant cells of step i) with said first compound X in a phytotoxic concentration and selecting plant cells comprising in their genome said first DNA construct, conferring resistance to said transformed plant cells against said compound X by expression of said D-amino acid oxidase, and
- 35 iv) providing at least one second compound M, which is non-phytotoxic or moderately phytotoxic against plant cells not functionally expressing said D-amino acid oxidase, wherein said compound M can be metabolized by said D-amino acid oxidase into one or more compound(s) N which are phytotoxic or more phytotoxic than compound M, and
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- 5 v) inducing deletion of said first expression cassette from the genome of said transformed plant cells and treating said plant cells with said second compound M in a concentration toxic to plant cells still comprising said first expression cassette, thereby selecting plant cells comprising said second expression cassette but lacking said first expression cassette.
4. The method of any of Claim 1 to 3 further comprising the step of regeneration of a fertile plant.
- 10 5. The method of any of Claim 1 to 4, wherein said first compound X is comprising a D-amino acid structure selected from the group consisting of D-tryptophane, D-histidine, D-arginine, D-threonine, D-methionine, D-serine, and D-alanine, and derivatives thereof.
- 15 6. The method of any of Claim 1 to 5, wherein said second compound M is comprising a D-amino acid structure selected from the group consisting of D-isoleucine, D-valine, D-asparagine, D-leucine, D-lysine, D-proline, and D-glutamine, and derivatives thereof.
- 20 7. The method of any of Claim 1 to 6, wherein deletion of said first expression cassette for the D-amino acid oxidase is realized by a method selected from:
- 25 a) recombination induced by a sequence specific recombinase, wherein said first expression cassette is flanked by corresponding recombination sites in a way that recombination between said flanking recombination sites results in deletion of the sequences in-between from the genome, or
- 30 b) homologous recombination between homology sequences A and A' flanking said first expression cassette, preferably induced by a sequence-specific double-strand break caused by a sequence specific endonuclease, wherein said homology sequences A and A' have sufficient length and homology in order to ensure homologous recombination between A and A', and having an orientation which – upon recombination between A and A' – will lead to excision of said first expression cassette from the genome of
- 35 said plant.
8. The method of Claim 7, wherein the recombinase or sequence-specific endonuclease, respectively, is expressed or combined with its corresponding recombination or recognition site, respectively, by a method selected from the group consisting of:
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- 5 a) incorporation of a second expression cassette for expression of the recombinase or sequence-specific endonuclease operably linked to a plant promoter into said DNA construct, preferably together with said first expression cassette flanked by said sequences which allow for specific deletion; or
- 10 b) incorporation of a second expression cassette for expression of the recombinase or sequence-specific endonuclease operably linked to a plant promoter into the plant cells or plants used as target material for the transformation thereby generating master cell lines or cells, or
- 15 c) incorporation of a second expression cassette for expression of the recombinase or sequence-specific endonuclease operably linked to a plant promoter into a separate DNA construct, which is transformed by way of co-transformation with said first DNA construct into said plant cells, or
- 20 d) incorporation of a second expression cassette for expression of the recombinase or sequence-specific endonuclease operably linked to a plant promoter into the plant cells or plants which are subsequently crossed with plants comprising the DNA construct of the invention.
- 25 9. The method of Claim 7 or 8, wherein deletion of said first expression cassette for the D-amino acid oxidase is induced or activated by inducing expression and/or activity of said sequence-specific recombinase or endonuclease by a method selected from the group consisting of
- 30 a) inducible expression by operably linking the sequence encoding said recombinase or endonuclease to an inducible promoter, and
- b) inducible activation, by employing a modified recombinase or endonuclease comprising a ligand-binding-domain, wherein activity of said modified recombinase or endonuclease can be modified by treatment of a compound having binding activity to said ligand-binding-domain.
- 35 10. The method of any of Claim 1 to 9, wherein the DNA construct is defined as in any of Claim 11 to 20 and the resulting plant cell or plant is selection marker free.
11. A DNA constructs suitable for the method of any of Claim 1 to 10, comprising

- 5 a) a first expression cassette comprising a nucleic acid sequence encoding a D-amino acid oxidase operably linked with a promoter allowing expression in plant cells or plants, wherein said first expression cassette is flanked by sequences which allow for specific deletion of said first expression cassette, and
- 10 b) at least one second expression cassette suitable for conferring to said plant an agronomically valuable trait, wherein said second expression cassette is not localized between said sequences which allow for specific deletion of said first expression cassette.
12. The DNA construct of Claim 11, wherein said D-amino acid oxidase expressed from said first expression cassette has preferably metabolizing activity against at least one D-amino acid and comprises the following consensus sequence:
- 15 [LVM]-[LVM]-H*-[NHA]-Y-G-x-[GSA]-[GSA]-x-G-x₅-G-x-A
- 20 wherein amino acid residues given in brackets represent alternative residues for the respective position, x represents any amino acid residue, and indices numbers indicate the respective number of consecutive amino acid residues.
- 25 13. The DNA construct of Claim 11 or 12, wherein said D-amino acid oxidase has enzymatic activity against at least one of the amino acids selected from the group consisting of D-alanine, D-serine, D-isoleucine, D-valine, and derivatives thereof.
- 30 14. The DNA construct of any of Claim 11 to 13 wherein said D-amino acid oxidase is described by a sequence of the group consisting of sequences described by GenBank or SwisProt Acc.No. JX0152, O01739, O33145, O35078, O45307, P00371, P14920, P18894, P22942, P24552, P31228, P80324, Q19564, Q28382, Q7PWX4, Q7PWY8, Q7Q7G4, Q7SFW4, Q7Z312, Q82MI8, Q86JV2, Q8N552, Q8P4M9, Q8PG95, Q8R2R2, Q8SZN5, Q8VCW7, Q921M5, Q922Z0, Q95XG9, Q99042, Q99489, Q9C1L2, Q9JXF8, Q9V5P1, Q9VM80, Q9X7P6, Q9Y7N4, Q9Z1M5, Q9Z302, and U60066.
- 35 15. The DNA construct of any of Claim 11 to 14 wherein said D-amino acid oxidase is selected from the group of amino acid sequences consisting of
- 40 a) the sequences described by SEQ ID NO: 2, 4, 6, 8, 10, 12, and 14, and
- b) the sequences having a sequence homology of at least 40% with a sequence as described by SEQ ID NO: 2, 4, 6, 8, 10, 12, and 14, and

- c) the sequences hybridizing under low or high stringency conditions with a sequence as described by SEQ ID NO: 2, 4, 6, 8, 10, 12, and 14.
- 5 16. The DNA construct of any of Claim 11 to 15, wherein said sequences which allow for specific deletion of said first expression cassette are selected from the group of sequences consisting of
- 10 a) recombination sites for a sequences-specific recombinase arranged in a way that recombination between said flanking recombination sites results in deletion of the sequences in-between from the genome, and
- 15 b) homology sequences A and A' having a sufficient length and homology in order to ensure homologous recombination between A and A', and having an orientation which – upon recombination between A and A' – will result in deletion of the sequences in-between from the genome.
- 20 17. The DNA construct of Claim 16, wherein said recombination sites correspond to a recombinase selected from the group consisting of the cre recombinase, the FLP recombinase, the Gin recombinase, the Pin recombinase, and the R recombinase.
- 25 18. The DNA construct of Claim 16, wherein said DNA construct comprises a recognition site of at least 10 base pairs for a sequence specific endonuclease between said homology sequences A and A'.
- 30 19. The DNA construct of Claim 18, wherein said recognition site corresponds to a sequence-specific endonuclease selected from the group consisting of the homing endonucleases I-SceI, I-CpaI, I-CpaII, I-CreI, and I-ChuI and chimeras thereof with ligand-binding domains.
- 35 20. The DNA construct of any of Claim 16 to 19, wherein said DNA construct further comprises an expression cassette for the sequence specific endonuclease or recombinase suitable for mediating deletion of the first expression cassette for the D-amino acid oxidase.
- 40 21. The DNA construct of Claim 20, wherein expression and/or activity of said sequence-specific recombinase or endonuclease can be induced and/or activated by a method selected from the group consisting of

- 5 a) inducible expression by operably linking the sequence encoding said recombinase or endonuclease to an inducible promoter,
- b) inducible activation, by employing a modified recombinase or endonuclease comprising a ligand-binding-domain, wherein activity of said modified re-combinase or endonuclease can be modified by treatment of a compound having binding activity to said ligand-binding-domain.
- 10 22. A transgenic vector comprising a DNA construct of any of Claim 11 to 21.
23. A transgenic cell comprising a DNA construct of any of Claim 11 to 21 or a vector of Claim 22.
24. A transgenic cell of Claim 23, wherein said cell is a plant cell.
- 15 25. A transgenic, non-human organism comprising a DNA construct of any of Claim 11 to 21, a vector of Claim 22, or a transgenic cell of Claim 23 or 24.
26. A transgenic, non-human organism of Claim 25 wherein said organism is a plant.
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